## ABSTRACT

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## 2,4-DISUBSTITUTED TRIAZINE DERIVATIVES

This invention concerns the use of the compounds of formula

$$\begin{array}{c|c}
 & R^{1} \\
 & A^{4} \\
 & A^{3} \\
 & A^{3}
\end{array}$$
(I)

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the *N*-oxides, the pharmaceutically acceptable addition salts, quaternary amines and the stereochemically isomeric forms thereof, wherein -a<sup>1</sup>=a<sup>2</sup>-a<sup>3</sup>=a<sup>4</sup>- forms a phenyl, pyridinyl, pyrimidinyl, pyridazinyl or pyrazinyl with the attached vinyl group; n is 0 to 4; and where possible 5; R<sup>1</sup> is hydrogen, aryl, formyl,  $C_{1.6}$ alkylcarbonyl,  $C_{1.6}$ alkyl,  $C_{1.6}$ alkyloxycarbonyl or substituted  $C_{1.6}$ alkyl; each R<sup>2</sup> independently is hydroxy, halo, optionally substituted  $C_{1.6}$ alkyl,  $C_{2.6}$ alkenyl or  $C_{2.6}$ alkynyl,  $C_{3.7}$ cycloalkyl,  $C_{1.6}$ alkyloxy,  $C_{1.6}$ alkyloxycarbonyl, carboxyl, cyano, nitro, amino, mono- or di( $C_{1.6}$ alkyl)amino, polyhalomethyl, polyhalomethyloxy, polyhalomethylthio, - $S(=O)_pR^4$ , - $NH-S(=O)_pR^4$ , - $C(=O)R^4$ , -NH-C(=O)H, - $C(=O)NHNH_2$ ,- $NHC(=O)R^4$ ,- $C(=NH)R^4$  or a 5-membered heterocyclic ring; p is 1 or 2; L is optionally substituted  $C_{1.0}$ alkyl,  $C_{2.10}$ alkenyl,  $C_{2.10}$ alkynyl or  $C_{3.7}$ cycloalkyl; or L is -X- $R^3$  wherein  $R^3$  is optionally substituted phenyl, pyridinyl, pyrimidinyl, pyrazinyl or pyridazinyl; X is - $NR^1$ -, -NH-NH-, -N-N-, -O-, -C(=O)-, -CHOH-, -S-, -S(=O)- or -S(=O)<sub>2</sub>-; aryl is optionally substituted phenyl; for the manufacture of a medicine for the treatment of subjects suffering from HIV (Human Immunodeficiency Virus) infection.